ESTHEMAX PHARM HAND SANITIZER- ethyl alcohol gel K Beauty Inc

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

esthemax pharm hand sanitizer

Active Ingredient

Ethyl Alcohol 83.0 %

Purpose

Antimicrobial

Warnings

Warnings

Flammable. Keep away from fire or flame. For external use only.

When using this product do not use in or near the eyes.

In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash appears and lasts.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Keep out of reach of children.

Directions

Place enough product in the palm of your hands to thoroughly cover your hands. Rub hands together briskly until product is completely absorbed and hands are dry.

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Inactive Ingredients:

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Inactive Ingredient: water

HAND SANITIZER SPRAY Kills 99.9% of Germs, Bacteria & Virus





Drug Facts

Active Ingredient

Purpose

Ethyl Alcohol 83% v/v...

.Antimicrobial

Uses • for hand rub to decrease bacteria on the skin • recommended for repeated use

Warnings

For external use only.

Flammable, keep away from heat and flame

Do not use in the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation and redness develop and persist for more than 72 hours.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately.

Directions: • wet hands thoroughly with product • briskly rubs hands together until dry • supervise children under 6 years in the use of this product.

Other information

- store at 68F to 77F
- · may discolor certain fabrics

Inactive Ingredient: Pure Water.

Distributed by esthemax

3555 W Lomita Blvd. Unit J Torrance, CA 90505

www.esthemaxpharm.com

MADE IN KOREA

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ESTHEMAX PHARM HAND SANITIZER

ethyl alcohol gel

Product Information

esthemax+pharm

3.5FL,OZ (100ml)

Product Type HUMAN OTC DRUG Item Code (Source) NDC:74926-0010

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name
Basis of Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)
ALCOHOL
83 mL in 100 mL

Inactive Ingredients

Ingredient Name Strength
GLYCERIN (UNII: PDC6A3C0OX)
WATER (UNII: 059QF0KOOR)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:74926- 0010-1	100 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/08/2020			
Marketing Information						
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		

04/08/2020

Labeler - K Beauty Inc (057989751)

part333A

OTC monograph not final

Establishment						
Name	Address	ID/FEI	Business Operations			
GL Pharm Co., Ltd.		694506955	manufacture(74926-0010)			

Revised: 3/2021 K Beauty Inc